

# EXHIBIT 1

**Declaration of Michael D. West, PhD  
CEO, LifeCraft Sciences, Inc.**

My name is Dr. Michael David West. I am a cell and molecular biologist with over 40 years of experience in biological and medical research relating to the biology of embryonic development and aging. I received my PhD at Baylor College of Medicine in the division of Molecular Virology where I was formally trained in virology and epidemiology. While I have participated in academic life as an adjunct Professor at the University of California, Berkeley, the majority of my career has been in the field of drug development in the biotechnology industry. I have founded or managed as CEO five public companies and led the development of numerous therapeutic products including some still in clinical trials and one currently FDA-approved and one marketed under the name RYTELO, for the treatment of myelodysplastic syndrome. I am an author on over 70 peer-reviewed scientific publications, an inventor on over 400 patents and patent applications filed world-wide, and have testified numerous times before the U.S. Congress as an expert in the field of embryonic development and cell-based regenerative therapies.

In preparation of this declaration, I reviewed the Curriculum Vitae of Ms. Kseniia Petrova as well as her peer-reviewed scientific publications. In addition, I carefully reviewed certain protocols used to prepare tissue samples of developing frog (*Xenopus*) embryos that Ms. Petrova was bringing to Harvard Medical School in February of 2025. To be certain that I was reviewing correct information, I emailed Dr. Anne-Hélène Monsoro-Burq of Institut Curie on Saturday May 24, 2025 and just received a detailed reply today (May 27, 2025).

Based on my training, knowledge, and expertise, in virology and epidemiology, cell biology, and managing pathogen-free containment for human clinical studies, I can affirm with certainty that the complete assortment of samples Ms. Petrova was transporting in February of this year was entirely non-hazardous, noninfectious, non-toxic, non-human histological samples as stated in the letter dated February 20, 2025 by Dr. Geneviève Almouzni. These developing frog samples that were being transported by Ms. Petrova to

Harvard Medical School were, in my opinion, entirely safe and posing no health safety risk of any type to humans, animals, agriculture, or any other known risk to the United States.

The reason this can be stated with certainty is that there is a plethora of scientific studies demonstrating that the formalin/formaldehyde treatments, together with alcohol dehydration render bacteria, bacterial spores, viruses, or fungi non-viable. By way of nonlimiting example, even relatively resistant bacterial spores are rendered inert by formalin treatment (Vardaxis et al, 1997. Sporicidal activity of chemical and physical tissue fixation methods. *J. Clin. Pathol.* 50: 429-433). All of the samples transported by Ms Petrova in February of 2025 were, according to representations made to me in an email from Dr. Anne-Hélène Monsoro-Burq of Institut Curie, treated in this manner. This is to be expected since this is standard protocol for this type of research. Therefore, put simply, the samples, while derived from living cells, were at the time of transport entirely inert in nature, similar to the way a piece of paper, while derived from plant material, should not be considered a “plant” in a biological context. Similarly, shoe leather, although biological in origin, by the process of tanning is rendered non-viable and paper and leather are not believed to be a possible vector for the transmission of pathogens.

Secondly, I can affirm that Ms. Kseniia Petrova is a scientist with highly-desirable skills for the United States in computer science as it relates to medical research, a field known as bioinformatics. I base this on her Curriculum Vitae and after carefully reviewing her peer-reviewed scientific publications. For example, she is the first author of a paper published in the prestigious scientific journal *Developmental Biology* titled “A Revised Single-Cell Transcriptomic Atlas of *Xenopus* Embryo Reveals New Differentiation Dynamics,” (*Dev Biol* 2024. 511: 76-83). Being first author means that she played the lead role in the paper. In this report, she describes the mapping of molecular pathways that could play a fundamental role in understanding the process of development and tissue regeneration, and novel therapies to address one of the largest challenges facing the United States;

namely, that of chronic age-related degenerative diseases such as heart failure, stroke, arthritis, as well as numerous other devastating human medical conditions.

The laboratory model of *Xenopus* is a well-established model of development and regeneration research. I am the founder of an electronic database on such pathways produced in collaboration with scientists in Israel called LifeMap Discovery where \$7M was expended to begin a mapping project in mice similar to that being undertaken by Ms. Petrova. If *Xenopus* development and regeneration could be so mapped, it could greatly advance medical research and likely, in my opinion, lead to therapies that could be of enormous benefit to citizens of the United States.

In summary, Ms. Petrova has skills that can easily be identified by researchers in the biological and medical sciences as being of enormous benefit to the United States if it is to continue to lead in the medical sciences. And her transport of fixed, denatured, and inert *Xenopus*-derived samples posed no conceivable health or safety risk to the United States and should not have been considered “biologicals” in the sense that they were alive or were a vector for a living pathogen.

I declare under penalty of perjury under the laws of the United States of America that the foregoing is true and correct to the best of knowledge and belief.

May 27, 2025

A handwritten signature in blue ink, appearing to read 'MD West', is written above a horizontal line.

Michael D. West, PhD  
CEO  
LifeCraft Sciences, Inc.

**MICHAEL DAVID WEST, B.S., M.S., Ph.D.****CURRICULUM VITAE****May, 2025****Summary:**

Dr. West is the CEO of LifeCraft Sciences, Inc. He has focused his career primarily on developing therapeutics targeting human aging. More specifically, his interests are the overlap of aging and regeneration and methods of reprogramming the cells of the human body back to a regenerative state. He received his Ph.D. from Baylor College of Medicine in 1989 concentrating on the biology of cell senescence and the secretome of aged cells (a phenomenon that came to be known as “SASP”). His experience includes being founder/CEO of five publicly-traded companies including being the founder and first CEO of Geron Corporation of Menlo Park, California (Nasdaq: GERN) where he was variously, CEO, Director, and Vice President and initiated and managed programs in telomerase diagnostics, oligonucleotide-based telomerase inhibition as anti-tumor therapy, and the cloning and use of the catalytic component of telomerase (*TERT*) in therapy to rescue cells from senescence. His pioneering work in telomerase inhibition led to the drug RyTelo® (Imetelstat) for the treatment of cancer. From 1995 to 1998 while at Geron he organized and managed the research collaboration between the company and its academic collaborators James Thomson and John Gearhart that led to the first isolation of human embryonic stem and human embryonic germ cells. From 1998 to 2007 he held positions as CEO, President, and Chief Scientific Officer at Advanced Cell Technology, Inc (later renamed Ocata), which was acquired by Astellas Pharma, Inc. At ACT he managed programs in reprogramming developmental aging using somatic cell nuclear transfer, retinal differentiation, and *PureStem*<sup>TM</sup>, a technology for the multiplex derivation and characterization of diverse clonal human embryonic progenitor cell lines. From 2007 to 2018 he was a CEO of Lineage Cell Therapeutics (NYSE American: LCTX) (previously known as BioTime, Inc.) where he led efforts to acquire Geron’s stem cell assets initially residing in Asterias Biotherapeutics (NYSE American: AST). During his tenure at Lineage, he led in the development of *OpRegen* for age-related macular degeneration which was eventually licensed to Genentech/Roche. He also led the development of *Renevia*, a matrix for cell transplantation now approved for use in Europe, *OPC-1* for the treatment of spinal cord injury, *VAC-2*, a human ES cell-derived dendritic cell vaccine targeting telomerase in lung cancer, and diverse novel preclinical programs in orthopedics and other applications in regenerative medicine. He founded AgeX Therapeutics (NYSE American: AGE) later merged with Serena Therapeutics (NYSE American: SER), where he led preclinical-stage programs in metabolic and vascular aging as well as technologies for the reversal of the aging of human cells to a youthful regenerative state (induced Tissue Regeneration (iTR)) commonly referred to as “partial reprogramming.” He also led allied programs related to cancer. He currently is founder and CEO of LifeCraft Sciences, Inc managing novel preclinical programs targeting aging and cancer.

Dr. West is an author of over 70 peer-reviewed scientific papers with over 3,300 citations, an inventor on over 370 patents and patent applications filed world-wide, lectures widely in scientific and pharmaceutical meetings, and has represented the scientific research and biotechnology industry in numerous Congressional Hearings. He has appeared on numerous television and radio programs, and podcasts including Meet the Press, CNN, CNBC, Fox News, The Oprah Winfrey Show, The Royal Society of London, BBC, Bill Nye, as well as others.

**Personal:**

Date and Place of Birth	April 28, 1953; Niles, Michigan
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**Education:**

Niles Senior High School; Niles, MI		1971
Rensselaer Polytechnic Institute; Troy, NY	B.S. Major: Psychology Minor: Management	1976
Andrews University; Berrien Springs, MI	M.S. Biology	1982
Baylor College of Medicine; Houston, TX	Ph.D. Cell Biology (Division of Molecular Virology)	1989
University of Texas Southwestern Medical Center at Dallas; Dallas, TX	Postdoc and Med. Student (MSIII)	1988-1992

**Business Experience:**

Geron Corporation (Nasdaq: GERN)	Founder, Director, Officer	1990-1998
Origen Therapeutics, Inc.	Founder, Chairman,	1997-1999
Advanced Cell Technology, Inc. (AKA Ocata Therapeutics, now now Astellas Pharma, Inc.)	Chairman & CEO	1998-2004
(NASDAQ: OCAT)	CSO	2004-2007
Cyagra, LLC.	President & CEO	2000-2002
A.C.T. Group, Inc.	Founder, Chairman, President & CEO	1999-2005
BioTime, Inc. (NYSE MKT: BTX, now LCTX) 2007-2015	CEO	
	Co-CEO	2015-2018
ReCyte Therapeutics	CEO	2007-2023
OncoCyte Corporation (NYSE: OCX)	Founder, Board Member	2009-2016
BioTime Asia	CEO	2010-2018
ES Cell International	CEO	2010-2013
OrthoCyte Corporation	CEO	2010-2018
LifeMap Sciences (GeneCards)	Board Member/CSO	2010-2021
Asterias Biotherapeutics (NYSE MKT: AST, now LCTX)	Board Member, VP	2013-2018
Asterias Biotherapeutics (NYSE MKT: AST, now LCTX)	CEO	2014
AgeX Therapeutics (NYSE: AGE now SER)	CEO	2017-2023
Reverse Bioengineering, Inc.	Founder, President	2019-2023
LifeCraft Sciences, Inc.	Founder, CEO	2024-Present

**Additional Affiliations:**

Ben-Gurion University of the Negev – Board of Governors

**Research and Professional Experience:**

2024-Present	LifeCraft Sciences, Inc. Alameda, CA 94501
2019-2023	Reverse Bioengineering, Inc. Founder, President 1101 Marina Village Parkway Alameda, CA 94501

2017-2023	Founder, Chief Executive Officer AgeX Therapeutics, Inc. (NYSE American: AGE, now SER) 1101 Marina Village Parkway Alameda, CA 94501
2010-2018	Chief Executive Officer OrthoCyte Corporation 1301 Harbor Bay Pkwy Alameda, CA 94502
2014-2014	Interim Chief Executive Officer Asterias Biotherapeutics (NYSE American AST, now LCTX) 230 Constitution Menlo Park, CA
2010-2013	Chief Executive Officer ES Cell International Pte. Ltd. Biopolis Street #01-03 Genome Singapore 138672
2010-2013	Chief Executive Officer BioTime Asia Hong Kong
2009-2011	Chief Executive Officer OncoCyte Corporation 1301 Harbor Bay Pkwy Alameda, CA 94502
2015-2018	Co-Chief Executive Officer BioTime, Inc. (NYSE American: BTX, now LCTX)) 1010 Atlantic Ave. Alameda, CA 94501
2007-2015	Chief Executive Officer BioTime, Inc. (NYSE American: BTX, now LCTX) 1010 Atlantic Ave. Alameda, CA 94501
2007-Present	Chief Executive Officer ReCyte Therapeutics, Inc. 1301 Harbor Bay Pkwy Alameda, CA 94502



2005-2008	Adjunct Professor Department of Bioengineering University of California, Berkeley
2004-2007	President, Chief Scientific Officer Advanced Cell Technology, Inc. (Re-named Ocata: Nasdaq: OCAT, now Astellas Pharma, Inc.) 381 Plantation St. Worcester, MA 01605
1998-2004	President, CEO, Chairman Advanced Cell Technology, Inc. (Re-named Ocata: Nasdaq: OCAT, now Astellas Pharma, Inc.) One Innovation Dr. Worcester, MA 01605
1999-2005	Founder, Chairman, President & CEO A.C.T. Group, Inc. (Holding company for OCAT) One Innovation Dr. Worcester, MA 01605
2003-Present:	Board Member BioTime, Inc.
2000-2002	President & CEO Cyagra, Inc. One Innovation Dr. Worcester, MA 01605
1997-1999	Founder, Chairman, Origen Therapeutics, Inc. 1450 Rollins Rd. Burlingame, CA 94080
1992-1998	Founder, VP, Chief Scientific Officer, VP New Technology Discovery (Nasdaq: GERN) Geron Corporation 230 Constitution Dr. Menlo Park, CA 94025
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## **RESEARCH GRANT SUPPORT**

2009 (PI: West); (06/09- 05/12); California Institute for Regenerative Medicine; Addressing the Cell Purity and Identity Bottleneck through Generation and Expansion of Clonal Human Embryonic Progenitor Cell Lines, \$4,721,706.

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3. West, M.D., Shay, J., Wright, W., Blackburn, E.H. 1993. Therapy and diagnosis of conditions related to telomere length and/or telomerase activity. PCT Publication No. WO1993023572A1.
4. Blackburn, E.H., Harley, C.B., Kim, N.W., Mceachern, M.J., Shay, J., Strahl, C.M., Vaziri, H., Weinrich, S.L., West, M.D., Wright, W.E. 1995. Therapy and Blackburn, E.H., Shay, J., West, M.D., Wright, W. 1993. Therapy and Diagnosis of Conditions Related to Telomere Length and/or Telomerase Activity. Australian Application No. AU4374093A.
5. West, M.D., Shay, J., Wright, W., Blackburn, E.H. 1993. Methods of modulating cell proliferation through telomerase modulation. Canadian Application No. CA2245462A1.
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11. West, M.D. 1996. Trypsin-sensitive Agent Able to Reduce PAI-1 and beta-APP Expression in Senescent Cells and Increase the Ability of Fibroblasts to Divide in Culture. U.S. Patent No. 5,482,838.
12. West, M.D. 1996. Isoquinoline poly (ADP-ribose) polymerase inhibitors to treat skin diseases associated with cellular senescence. U.S. Patent No. 5,589,483.
13. West, M.D., Shay, J., Wright, W. 1996. Therapy and Diagnosis of Conditions Related to Telomere Length and/or Telomerase Activity. U.S. Patent No. 5,489,508.
14. West, M., Shay, J., Wright, W., Blackburn, E. 1996. Treatment and diagnosis of conditions associated with telomere length and / or telomerase activity. Japanese Application No. JPH08501079A.
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33. Bodnar, A., Chiu, C-P, Gold, J.D., Inokuma, M., Murai, J.T., West, M.D. 1999. Methods and materials for the growth of primate-derived primordial stem cells in feeder-free culture. Canadian Application No. CA2307807A1.

34. Bodnar, A., Chiu, C-P, Gold, J.D., Inokuma, M., Murai, J.T., West, M.D. 1999. Methods and materials for the growth of primate-derived primordial stem cells. PCT Publication No. WO1999020741A1.
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58. West, M.D. 2001. The production of agricultural animals from embryonic stem (es) cells. Australian Application No. AU5970301A.
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